

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application.

Listing of Claims:

Claims 1 – 5. Cancelled.

Claims 6 – 8. Cancelled.

Claim 9 (Currently Amended). A therapeutic vaccine against tuberculosis comprising one or more polypeptides or fragments thereof, which polypeptides are upregulated or expressed during the latent stage of the mycobacteria infection, according to claim 28, wherein at least one of the one or more polypeptides or fragments thereof, is fused to at least one fusion partner which is an antigen expressed by bacteria within the mycobacteria family.

10 (Previously Presented). A therapeutic vaccine according to claim 9 where the fusion partner is selected from the group consisting of ESAT-6, ESAT-6-Ag85B, TB10.4, CFP10, RD1-ORF5, RD1-ORF2, Rv1036, MPB64, MPT64, Ag85A, Ag85B (MPT59), MPB59, Ag85C, 19kDa lipoprotein, MPT32.

Claim 11. Cancelled.

12 (Currently Amended). A multiphase vaccine according to claim 28, further comprising antigen components with therapeutic activity against tuberculosis comprising one or more polypeptides or fragments thereof, which polypeptides are upregulated or expressed during the latent stage of the mycobacteria infection, combined with antigen components with prophylactic activity.

13 (Currently Amended). A ~~multi~~phase vaccine according to claim 12 where the antigen components with prophylactic activity are selected from the group consisting of ESAT-6, ESAT-6-Ag85B, TB10.4, CFP10, RD1-ORF5, RD1-ORF2, Rv1036, MPB64, MPT64, Ag85A, Ag85B (MPT59), MPB59, Ag85C, 19kDa lipoprotein or MPT32.

Claims 14 and 15. Cancelled.

Claim 16 (Currently Amended). A therapeutic vaccine ~~against tuberculosis~~ comprising ~~one or more polypeptides or fragments thereof and a pharmaceutically acceptable carrier, vehicle or adjuvant, which polypeptides are upregulated or expressed during the latent stage of the mycobacteria infection according to claim 28,~~ where the polypeptides or fragments are recombinant or synthetic and are delivered in a delivery system such as an adjuvant.

Claim 17 (Currently Amended). A therapeutic vaccine ~~against tuberculosis~~ comprising ~~one or more polypeptides or fragments thereof and a pharmaceutically acceptable carrier, vehicle or adjuvant, which polypeptides are upregulated or expressed during the latent stage of the mycobacteria infection according to claim 28,~~ in which the polypeptide or fragment is lipidated so as to allow a self-adjuvanting effect of the polypeptide.

Claims 18 – 23. Cancelled.

24 (Currently Amended). The therapeutic vaccine according to claim 28 6, wherein the polymeric carrier and the one or more polypeptide(s) are bound by non-covalent interactions.

25 (Previously Presented). The therapeutic vaccine according to claim 24, wherein the polymeric carrier is a polystyrene.

26 (Currently Amended). The therapeutic vaccine according to claim 28 6, wherein the polymeric carrier and the one or more polypeptide(s) or fragments thereof are bound by covalent interactions.

27 (Previously Presented). The therapeutic vaccine according to claim 26, wherein the polymeric carrier is selected from the group consisting of a polysaccharide or a polypeptide.

28 (Previously Presented). A therapeutic vaccine against tuberculosis comprising one or more mycobacteria polypeptides and a pharmaceutically acceptable polymeric carrier bound to the one or more polypeptides or a pharmaceutically acceptable adjuvant, which polypeptides are upregulated or expressed during the latent stage of the mycobacteria infection which is characterized by low-oxygen tension in the microenvironment of the mycobacteria.

29 (Currently Amended) The therapeutic vaccine according to claim 28 4, wherein the one or more polypeptides has an amino acid sequence selected from SEQ ID NO 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, 12, 13, 14, 15, 16, 17, 18, 19, 20, 21, 22, 23, 24, 25, 26, 27, 28, 29, 30, 31, 32, 33, 34, 35, 36, 37, 38, 39, 40, 41, 42, 43, 44 and 45.